AMENDMENTS TO THE CLAIMS

- 1. (Currently Amended) A packaged tablet, which tablet has a matrix consisting of comprising at least 55% of a cellulose ether, wherein the tablet has a water activity of at most 0.6 and is packaged such as to delay moisture uptake by the tablet.
- 2. (Previoulsy Presented) The packaged tablet according to claim 1, wherein the tablet has a water activity of less than 0.55.
- 3. (Currently Amended) A packaged tablet, which tablet has a matrix eonsisting of comprising at least 55% of a cellulose ether, wherein the tablet has a water content of less than 9% w/w and is packaged such as to delay moisture uptake by the tablet.
- 4. (Currently Amended) The packaged tablet according to claim 1, wherein the matrix eonsists of comprises more than 65% of a cellulose ether.
- 5. (Previoulsy Presented) The packaged tablet according to claim 4, wherein the cellulose ether is hydroxypropyl methylcellulose.
- 6. (Currently Amended) The packaged tablet according to claim 1, wherein the tablet comprises gepirone HCl in an amount in the range of from 20-85 mg.
- 7. (Currently Amended) The packaged tablet according to claim 3, wherein the matrix eonsists of comprises more than 65% of a cellulose ether.

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- 8. (Currently Amended) The packaged tablet according to claim 3, wherein the tablet comprises gepirone HCl in an amount in the range of from 20-85 ma 20-85 mg.
- 9. (Currently Amended) The packaged tablet according to claim 1, wherein the tablet comprises gepirone HCl in an amount in the range of from 20-85 ma 20-85 mg and the matrix consists of comprises more than 65% of a cellulose ether.
- 10. (Previoulsy Presented) The packaged tablet according to claim 9, wherein the cellulose ether is hydroxypropyl methylcellulose.
- 11. (New) The packaged tablet according to claim 1, wherein the matrix comprises 70 to 85% of a cellulose ether.
- 12. (New) The packaged tablet according to claim 1, wherein the tablet comprises gepirone HCl in an amount in the range of from 60-85 mg.
- 13. (New) The packaged tablet according to claim 1, wherein the matrix further comprises microcrystalline cellulose.
- 14. (New) The packaged tablet according to claim 1, wherein the matrix further comprises euroxide, yellow ferric oxide, and/or red ferric oxide.
- 15. (New) The packaged tablet according to claim 1, wherein the matrix further comprises colloidal anhydrous silicon dioxide.

- 16. (New) The packaged tablet according to claim 1, wherein the matrix further comprises magnesium stearate.
- 17. (New) The packaged tablet according to claim 3, wherein the matrix comprises 70 to 85% of a cellulose ether.
- 18. (New) The packaged tablet according to claim 3, wherein the tablet comprises gepirone HCl in an amount in the range of from 60-85 mg.
- 19. (New) The packaged tablet according to claim 3, wherein the matrix further comprises microcrystalline cellulose.
- 20. (New) The packaged tablet according to claim 3, wherein the matrix further comprises euroxide, yellow ferric oxide, and/or red ferric oxide.
- 21. (New) The packaged tablet according to claim 3, wherein the matrix further comprises colloidal anhydrous silicon dioxide.
- 22. (New) The packaged tablet according to claim 3, wherein the matrix further comprises magnesium stearate.

SUPPORT FOR THE AMENDMENTS

Claims 1, 3, 4, and 6-9 have been amended.

Claims 11-22 have been added.

The amendment of Claims 1, 3, 4, and 6-9 and new Claims 11-22 is supported by the originally filed claims and by the specification at pages 2, line 29 to page 4, line 30 and Table 1 on page 5.

No new matter has been entered by the present amendment.

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